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The listing of claims will replace all prior versions, and listings, of claims in the application: Listing of Claims:

1. (Currently Amended) A compound of formula (II), or a pharmaceutical or veterinarily acceptable salt, hydrate or solvate thereof

$$Q \xrightarrow{R_1} Q \xrightarrow{R_3} Q \xrightarrow{R_4} Q \xrightarrow{N} Q \xrightarrow{R_5} (II)$$

wherein

Q represents a radical of formula -N(OH)CH(=O) or formula -C(=O)NH(OH);

R₁ represents hydrogen, methyl or trifluoromethyl, or, except when Z is a radical of formula -N(OH)CH(=O), a hydroxy, halo or amino group;

 R_2 represents a group R_{10} - $(V)_n$ - $(ALK)_m$ - wherein

R₁₀ represents hydrogen, or a C₁-C₆ alkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, cycloalkyl, aryl, or heterocyclyl group, any of which may be unsubstituted or substituted by (C₁-C₆)alkyl, (C₁-C₆)alkoxy, hydroxy, mercapto, (C₁-C₆)alkylthio, amino, halo (including fluoro, ehloro, bromo and iodo), trifluoromethyl, cyano, nitro, oxo, -COOH, -CONH₂, -COOR^A, -NHCOR^A, -CONHR^A, -NHR^A, -NR^AR^B, or -CONR^AR^B wherein R^A and R^B are independently a (C₁-C₆)alkyl group and

ALK represents a straight or branched divalent C₁-C₆ alkylene, C₂-C₆ alkenylene, or C₂-C₆ alkynylene radical, and may be interrupted by one or more non-adjacent -NH-, -O- or -S-linkages,

V represents -NH-, -O- or -S-, and

m and n are independently 0 or 1;

R₃ represents the side chain of a natural or non-natural alpha amino acid; R₄ represents hydrogen or C₁-C₃ alkyl;

Y represents N or CH;

ring A is optionally substituted on one or more ring carbon atoms by C_1 - C_3 alkyl, C_1 - C_3 alkoxy, or halo; and

R₅ represents a group (IIA),

$$\rightarrow$$
 (IIA)

wherein

m is 0 or 1;

Alk¹ represents a divalent C₁-C₃ alkylene radical;

Z represents hydrogen or cycloalkyl, phenyl or heterocyclic which is optionally substituted by

(C₁-C₆)alkyl,

phenyl,

monocyclic 5 or 6-membered heterocyclic,

benzyl,

phenoxy, or (C₁-C₆)alkoxy,

phenylthio or (C₁-C₆)alkylthio, any of which is in turn optionally substituted by:

hydroxy or mercapto,

trifluoromethyl,

oxo,

nitro,

cyano (-CN),

bromo, chloro, fluoro, or iodo,

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-COOH, or -COOR^A,
-CONH_2, -CONHR^A, or -CONR^AR^B
-COR^A, -SO_2R^A,
-NHCOR^A,
-NH_2, -NHR^A, or -NR^AR^B,

wherein R^A and R^B are independently a (C_1-C_6)alkyl group, or R^A
and R^B taken together with the nitrogen atom to which they are attached form a 5- or 6-membered heterocyclic ring which may be substituted by-(C_4C_3)alkyl, hydroxy, or hydroxy(C_1-C_3)alkyl.
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2. (Currently Amended) A compound as claimed in claim 1 wherein Z represents cycloalkyl, phenyl or monocyclic-heterocyclic, which is optionally substituted by

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(C<sub>1</sub>C<sub>6</sub>)alkyl, (C<sub>2</sub>-C<sub>6</sub>)alkenyl, or (C<sub>2</sub>-C<sub>6</sub>)alkynyl,
phenyl, or halophenyl,
trifluoromethyl,
monocyclic 5 or 6-memberedhetrocyclic,
benzyl, or halophenylmethyl,
hydroxy, phenoxy, (C<sub>1</sub>-C<sub>6</sub>)alkoxy, or hydroxy(C<sub>1</sub>-C<sub>6</sub>)alkyl,
mercapto, (C<sub>1</sub>-C<sub>6</sub>)alkylthio or mercapto(C<sub>1</sub>-C<sub>6</sub>)alkyl,
oxo,
nitro,
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cyano (-CN),

bromo, chloro, fluoro, or iodo,

-COOH, or -COOR^A,

-CONH₂, -CONHR^A, or-CONR^AR^B

 $-COR^A$, $-SO_2R^A$,

-NHCOR^A,

-NH₂, -NHR^A, or -NR^AR^B,

wherein R^A and R^B are independently a $(C_1\text{-}C_6)$ alkyl group, or R^A and R^B taken together with the nitrogen atom to which they are attached form a 5-or 6-membered heterocyclic ring which may be substituted by $(C_1\text{-}C_3)$ alkyl, hydroxy, or hydroxy $(C_1\text{-}C_3)$ alkyl.

- 3. (Currently amended) A compound as claimed in claim $1 \frac{1}{\text{or elaim 2}}$ -wherein R_1 is hydrogen.
- 4. (Currently amended) A compound as claimed in any of the preceding claims claim 1 wherein R_2 is (C_1-C_6) alkyl-, cycloalkyl (C_1-C_6) alkyl-, (C_1-C_3) alkyl-S- (C_1-C_3) alkyl-, or (C_1-C_3) alkyl-O- (C_1-C_3) alkyl-.
- 5. (Currently amended) A compound as claimed in any of claims 1 to 3 claim 1 wherein R₂ is n-propyl, n-butyl, n-pentyl, cyclopentylmethyl, cyclopentylethyl, cyclohexylmethyl or cyclohexylethyl.
- 6. (Currently amended) A compound as claimed in any of the preceding claims claim 1

the characterising group of a natural α amino acid, for example benzyl, or 4methoxyphenylmethyl, in which any functional group may be protected, any amino group may be acylated and any carboxyl group present may be amidated; or a group - $[Alk]_nR_9$ where Alk is a (C_1-C_6) alkylene or (C_2-C_6) alkenylene group optionally interrupted by one or more -O-, or -S- atoms or -N(R_{12})- groups [where R_{12} is a hydrogen atom or a (C_1-C_6) alkyl group], n is 0 or 1, and R_9 is hydrogen or an optionally substituted phenyl, aryl, heterocyclyl, cycloalkyl or cycloalkenyl group or (only when n is 1) R₉ may additionally be hydroxy, mercapto, (C₁-C₆)alkylthio, amino, halo, trifluoromethyl, nitro, -COOH, -CONH₂, -COOR^A, -NHCOR^A, -CONHR^A, -NHR^A, -NR^AR^B, or -CONR^AR^B wherein R^A and R^B are independently a (C₁-C₆)alkylgroup; or a benzyl group substituted in the phenyl ring by a group of formula -OCH₂COR₈ where R_8 is hydroxyl, amino, (C_1-C_6) alkoxy, phenyl (C_1-C_6) alkoxy, (C_1-C_6) alkylamino, di $((C_1-C_6)$ alkoxy, phenyl (C_1-C_6) alkylamino, phenyl $(C_1-C$ C_6)alkyl)amino, phenyl(C_1 - C_6)alkylamino; or a heterocyclic (C₁-C₆)alkyl group, either being unsubstituted or mono- or disubstituted in the heterocyclic ring with halo, nitro, carboxy, (C_1-C_6) alkoxy, cyano, (C_1-C_6) alkanoyl, trifluoromethyl(C₁-C₆)alkyl, hydroxy, formyl, amino, (C₁-C₆)alkylamino, di-(C₁- C_6)alkylamino, mercapto, (C_1-C_6) alkylthio, hydroxy (C_1-C_6) alkyl, mercapto (C_1-C_6) alkyl or (C_1-C_6) alkylphenylmethyl; or a group-CR_aR_bR_c in which:

each of R_a, R_b and R_c is independently hydrogen, (C₁-C₆)alkyl, (C₂-C₆)alkenyl, (C_2-C_6) alkynyl, phenyl (C_1-C_6) alkyl, (C_3-C_8) cycloalkyl, or

R_c is hydrogen and R_a and R_b are independently phenyl or heteroaryl-such as

pyridyl; or

R_c is hydrogen, (C₁-C₆)alkyl, (C₂-C₆)alkenyl, (C₂-C₆)alkynyl, phenyl(C₁-C₆)alkyl, or (C₃-C₈)cycloalkyl, and R_a and R_b together with the carbon atom to which they are attached form a 3 to 8 membered cycloalkyl or a 5-to 6-membered heterocyclic ring; or

R_a, R_b and R_c together with the carbon atom to which they are attached form a tricyclic ring-(for example adamantyl); or

R_a and R_b are each independently (C₁-C₆)alkyl, (C₂-C₆)alkenyl, (C₂-C₆)alkynyl, phenyl(C₁-C₆)alkyl, or a group as defined for R_c below other than hydrogen, or R_a and R_b together with the carbon atom to which they are attached form a cycloalkyl or heterocyclic ring, and R is hydrogen, -OH, -SH, halogen, -CN, -CO₂H, (C₁-C₄) perfluoroalkyl, -CH₂OH, -CO₂(C₁-C₆)alkyl, -O(C₁-C₆)alkyl, -O(C₂-C₆)alkenyl, - $S(C_1-C_6)$ alkyl, $-SO(C_1-C_6)$ alkyl, $-SO_2(C_1-C_6)$ alkyl, $-S(C_2-C_6)$ alkenyl, $-SO(C_2-C_6)$ alkyl, $-SO(C_3-C_6)$ alkyl, $-SO(C_5-C_6)$ alkyl, $-SO(C_5-C_6)$ alkyl, $-SO(C_5-C_6)$ alkyl, $-SO(C_$ C₆)alkenyl, -SO₂(C₂-C₆)alkenyl or a group -Q-W wherein Q represents a bond or -O-, -S-, -SO- or -SO₂- and W represents a phenyl, phenylalkyl, (C_3-C_8) cycloalkyl, (C_3-C_8) cycloalkylalkyl, (C_4-C_8) cycloalkenyl, (C_4-C_8) cycloalkenylalkyl, heteroaryl or heteroarylalkyl group, which group W may optionally be substituted by one or more substituents independently selected from, hydroxyl, halogen, -CN, -CO₂H, -CO₂(C₁-C₆)alkyl, -CONH₂, -CONH(C₁-C₆)alkyl alkyl, -CONH(C₁-C₆alkyl)₂, -CHO, -CH₂OH, (C₁-C₄)perfluoroalkyl, - $O(C_1-C_6)$ alkyl, $-S(C_1-C_6)$ alkyl, $-SO(C_1-C_6)$ alkyl, $-SO_2(C_1-C_6)$ alkyl, $-NO_2$, $-NH_2$, - $NH(C_1-C_6)alkyl$, $-N((C_1-C_6)alkyl)_2$, $-NHCO(C_1-C_6)alkyl$, $(C_1-C_6)alkyl$, $(C_2-C_6)alkyl$, $(C_1-C_6)alkyl$, $(C_1-C_6)alkyl$, $(C_1-C_6)alkyl$, $(C_2-C_6)alkyl$, $(C_1-C_6)alkyl$ C₆)alkenyl, (C₂-C₆)alkynyl, (C₃-C₈)cycloalkyl, (C₄-C₈)cycloalkenyl, phenyl or

- 7. (Currently amended) A compound as claimed in any of claims 1 to 6 claim 1 wherein R₃ is methyl, ethyl, n-propyl, n-butyl, benzyl, 4-chlorobenzyl, 4-hydroxybenzyl, phenyl, cyclohexyl, cyclohexylmethyl, pyridin-3-ylmethyl, tert-butoxymethyl, naphthylmethyl, iso-butyl, sec-butyl, tert-butyl, 1-benzylthio-1-methylethyl, 1-methylthio-1-methylethyl, 1- mercapto-1-methylethyl, 1-methoxy-1-methylethyl, 1-hydroxy-1-methylethyl, 1-fluoro- 1-methylethyl, hydroxymethyl, 2-hydroxethyl, 2-carboxyethyl, 2-methylcarbamoylethyl, 2-carbamoylethyl, or 4-aminobutyl.
- 8. (Currently amended) A compound as claimed in any of claims 1 to 6 claim 1 wherein R₃ is tert-butyl, isobutyl, benzyl, isopropyl or methyl.
- 9. (Currently amended) A compound as claimed in any of the preceding claims claim 1 wherein R_4 is methyl.
- 10. (Currently amended) A compound, method, use or composition as claimed in any of the preceding claims as claimed in claim 1 wherein in the group R₅, m is 1, and Alk¹ is -(CH₂)- or (CH₂CH₂)-.
- 11. (Currently amended) A compound as claimed in any of the preceding claims claim 1 wherein, in the group R₅, Z is a phenyl, pyridyl, thienyl, furanyl, pyranyl, pyrolyl, diazolyl, triazolyl, thiazolyl, thiadiazolyl, oxazolyl, ozadiazolyl, indolyl, benzisozazolyl, benzthiazolyl or imidazothiazolyl ring, optionally substituted as specified in claim 1-of elaim 2.

- 12. (Original) A compound as claimed in claim 11 wherein the ring Z is unsubstituted or substituted by methyl, methoxy, ethoxy, methoxymethyl, ethylthio, chloro, bromo, hydroxy, nitro, phenyl, 2- or 4-nitrophenyl, dimethylamino, dimethylaminophenyl, methylsulphonyl, dimethylaminosulphonyl, 3-pyridyl or 2-pyrazin-2-yl.
- 12. (Canceled).
- 13. (Currently amended) A compound as claimed in claim 1or claim 2-wherein the compound is one specifically named and/or exemplified herein, or is the hydroxamate (Q represents a radical of formula -C(=O)NH(OH)) analogue thereof.
- 14. (Currently amended) A method for the treatment of bacterial infections in humans and non-human mammals, which comprises administering to a subject suffering such infection an antibacterially effective dose of a compound as claimed in any of claims 1 to 13 claim 1.
- 15. (Currently amended) A method for the treatment of bacterial contamination by applying an antibacterially effective amount of a compound as claimed in any of claims 1 to 13-claim 1 to the site of contamination.
- 16. (Currently amended) The use of a compound as claimed in any of claims 1 to 13 claim 1 in the manufacture of an antibacterial composition.

- 17. (Currently amended) A pharmaceutical or veterinary composition comprising a compound as claimed in any of claims 1 to 13-claim 1 together with a pharmaceutical of veterinarily acceptable carrier.
- 18. A compound as claimed in claim 1 wherein, in the group R₅, Z is a cyclopentyl, cyclohexyl, phenyl, morpholinyl, pyrimidin-2-yl, 1,2,3-thiadiazol-5-yl, 1,4-thiazol-5-yl, benzofuran-2-yl, 2- or 3-furanyl, 2- or 3-thienyl, 2- or 3-pyranyl, 2-, 3- or 4-pyrrolyl, 3-, 4- or 5-pyrazolyl, 3-, 4- or 5-isoxazolyl, or 2-, 3-or 4-pyridyl ring any of which may optionally be substituted by hydroxy, methoxy, ethoxy, mercapto, methylthio, ethylthio, methyl, ethyl, trfluoromethyl, fluoro, chloro, amino, methylamino, or dimethylamino.